UP - PGT CHEMISTRY

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(MOCK TEST)

- 1. An ideal gas undergoes isothermal expansion at constant pressure. During the process
 - (a) Enthalpy increases but entropy decreases
 - (b) Enthalpy remains constant but entropy increases
 - (c) Enthalpy decreases but entropy increases
 - (d) Both enthalpy and entropy remain constant.
- 2. A sealed container with gas at 2.00 atm is heated from 20.0 K to 40.0 K. The new pressure is:
 - (a) 0.50 atm
- (b) 1.00 atm
- (c) 4.00 atm
- (d) 2.14 atm
- 3. An element having an atomic radius of 0.14 nm crystallizes in an fcc unit cell. What is the length of a side of the cell?
 - (a) 0.96 nm
- (b) 0.4 nm
- (c) 0.24 nm
- (d) 0.56 nm
- 4. Which carbonyl compound has maximum dipole moment?

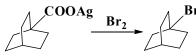








5. Following reaction goes through:



- (a) Carbene intermediate
- (b) Free radical intermediate
- (c) Carbocation intermediate
- (d) Carbanion intermediate
- 6. Oxidation product of quinoline with KMnO₄ is:
 - (a) Phthalic anhydride (b) Phthalic acid (c) Nicotinic acid (d) None of these
- 7. In the IR spectrum, carbonyl absorption band for the following compound appears at



- (a) 1810 cm⁻¹
- (b) 1770 cm⁻¹
- (c) 1730 cm⁻¹
- (d) 1690 cm⁻¹

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8. The compound that gives precipitate on warming with aqueous AgNO, is:









9. Which of the following alcohols would be most soluble in water?

(a) Propanol

(b) Hexanol

(c) Pentanol

(d) Butanol

10. Which of the reagent will give effective transformation of given compounds?

(a) CH₂N₂

(b) CH₂Li

(c) (CH₃),CuLi

(d) $Ph_{2}P = CH_{2}$

What is used to carry out the following conversion? 11.

- (a) Hydroboration oxidation followed by Jones oxidation
- (b) Wacker oxidation followed by haloform reaction
- (c) Oxymercuration demercuration followed by Jones oxidation
- (d) Ozonolysis followed by haloform reaction
- 12. What is a necessary condition for osmosis to take place?

(a) Semi–permeable membrane

(b) Same concentration of solvent

(c) High temperature

(d) Pressure greater than osmotic pressure

Which is the most appropriate method for determining the molar masses of 13. biomolecules?

(a) Relative lowering of vapor pressure (b) Elevation of boiling point

(c) Depression in freezing point

(d) Osmosis

14. If a reaction is of nth order the half life period of the initial concentration of reactants.

(a) Is independent

(b) Varies inversely as $(n-1)^{th}$ power

(c) Varies inversely at nth power

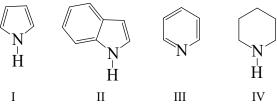
(d) Varies directly as $(n-1)^{th}$ power

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15.	The time taken for 10% completion of a first order reaction is 20 min. Then, for 19% completion, the reaction will take, given [log 81 = 1.908]								
	(a) 40 min	(b) 60 min	(c)) 30 min	(d) 50 min				
16.	The concept of t	is useful for	the reacti	on of:					
	(a) Zero order	(b) 1 order		2 order	(d) All orde	r			
17.	Which of the foll	owing represe	nt the corro	ect bond orde	rs for N ₂ , N ₂ + an	d N ₂ molecules?			
	(a) 3.0, 2.5, 2.5	(b) 3.0, 2.0	, 2.5 (c)	3.0, 3.0, 3.0	(d) 2.5, 2.5,	2.5			
18.	Which transition	ns are studies	by UV spec	ctrophotomer	.?				
	(a) Rotational	(b) Electron	nic (c)) Vibrational	(d) Nuclear				
19.	Which of the foll (a) The properties (b) 4f and 5f orbit (c) d-block eleme (d) 4d and 5d orbit	s of various act tals are equally ents show irreg	tinoids are v shielded gular and er	very similar	properties				
20.	Among the follo stable conforma		nds the co	mpounds hav	ing anti-confo	rmation as most			
	(a) F	(b) HO	OH (c) HO	F (q) Br	Br			
21.	The compound t	hat is antiaro	matic is						
		H B			0				
		I	II	III	IV				
	(a) I	(b) II	(c)) III	(d) IV				

- 22. A covalent molecule AB₃ has pyramidal structure. The number of lone pair and bond pair electrons in the molecule are respectively
 - (a) 2 and 2
- (b) 1 and 3
- (c) 0 and 4
- (d) 3 and 1
- 23. Complexes in which a metal is attached to only one kind of donor group is called
 - (a) Unidentate
- (b) Chelate
- (c) Homoleptic
- (d) Heteroleptic

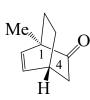
- 24. For the given complex [CoCl₂(en)(NH₃)₂]⁺, the number of geometrical isomers, the number of optical isomers and total number of isomers of all type possible respectively are
 - (a) 0, 1, 3
- (b) 0, 2, 2
- (c) 2, 2, 3
- (d) 3, 3, 4
- 25. The correct order of the basicity of the following compounds is



- (a) IV > III > II > I
- (c) IV > III > I > II

- (b) III > IV > II > I
- (d) III > IV > I > II
- 26. The donation of lone pair of electrons of CO carbon into the vacant orbital of metal atom results in bond.
 - (a) Sigma
- (b) pi
- (c) Back
- (d) Synergic
- 27. A complex having ______ geometry can have more than one type of hybridization.
 - (a) Tetrahedral

- (b) Square planar
- (c) Trigonal bipyramidal
- (d) Octahedral
- 28. The configuration at the two stereocentres in the compound given below are



- (a) 1R, 4R
- (b) 1R, 4S
- (c) 1S, 4R
- (d) 1S, 4S
- 29. In BrF₃ molecule the lone pairs occupy an equatorial position to minimize
 - (a) Lone pair-bond pair repulsion only
 - (b) Bond pair-bond pair repulsion only
 - (c) Lone pair-lone pair repulsion and lone pair-bond pair repulsion
 - (d) Lone pair—lone pair repulsion only

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30.	Identify	the	wrong	statement	in	the	following	ng

- (a) Atomic radius of the elements decreases as one move across from left to right in the 2nd period of the periodic table
- (b) Atomic radius of the elements increases as one move down the first group of the periodic table
- (c) Amongst isoelectronic species, greater the negative charge on the anion, larger is the ionic radius
- (d) Amongst isoelectronic species, smaller the positive charge on the cation, smaller is the ionic radius

31.	Theore	which	decreases	:4h	dilution	:.
31.	i ne one	willcii	uecreases	WILII	anunon	18

(a) Specific conductance

(b) Molar conductance

(c) Conductance

(d) Equivalent conductance

32. In which of the following octahedral complexes of Co (atomic no. 27), will the magnitude of Δ_{α} be the highest?

(a) $[Co(NH_3)_6]^{3+}$

(b) $[Co(CN)_{6}]^{3-}$ (c) $[Co(H_{2}O)_{6}]^{3+}$ (d) $[Co(C_{2}O_{4})_{3}]^{3-}$

33. On increasing the concentration of reactants in a reversible reaction, then equilibrium constant will

(a) Depend on the concentration

(b) Increase

(c) Unchanged

(d) Decrease

According to Wade's theory the anion [B₁,H₁,]²⁻ adopts 34.

(a) Closo-structure

(b) Arachno-structure

(c) Hypo-structure

(d) Nido-structure

35. The oxo-acid of phosphorus having P-atoms in +4, +3 and +4 oxidation states respectively, is

(a) $H_5P_3O_{10}$

(b) $H_5P_7O_7$

(c) $H_5P_2O_0$

 $(d) H_{5}P_{7}O_{0}$

With respect to enthalpy of combustion which of the following is correct? **36.**

(a) $\Delta H_{rxn} = \Sigma \Delta H_f(product) - \Sigma \Delta H_f(reactant)$

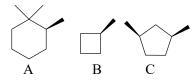
(b) $\Delta H_{rxn} = \Sigma \Delta H_f (reactant) - \Sigma \Delta H_f (product)$

(c) $\Delta H_{rxn} = \Sigma \Delta H_f (product) + \Sigma \Delta H_f (reactant)$

(d) $\Delta H_{rxn} = 2\Sigma\Delta H_f(product) - \Sigma\Delta H_f(reactant)$

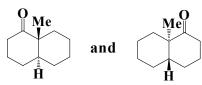
- 37. Which of the following molecules exists as a pair of enantiomers?
 - (a) 2–Bromopropane

- (b) 1-Bromo-3-methylbutane
- (c) 2–Cyclohexen–1–ol
- (d) cis-1,2-Dichlorocyclobutane
- 38. Which of the following compound(s) is/are chiral?



- (a) Only A and B
- (b) Only B
- (c) Only B and C
- (d) Only A
- **39.** How many chiral carbon atoms are present in 2, 3, 4-trichloropentane?
 - (a) 2
- (b) 1
- (c)3
- (d) 4

40. The two compounds given below are



- (a) Enantiomers
- (b) Identical
- (c) Diastereomers (d) Regioisomers
- 41. The molar volume of any ideal gas at N.T.P. is
 - (a) 10 litres
- (b) 44.8 litres
- (c) 21 litres
- (d) 22.4litres
- At what temperature are average speeds of CH₄. Will be same as that of O₂ at 300K 42.
 - (a) 150K
- (b) 300K
- (c) 600K
- (d) 900K

- The expected H H H bond angle in 43.
 - (a) 180°
- (b) 120°
- $(c) 60^{\circ}$
- (d) 90°
- 44. Which type of colloid is the dissolution of sulphur (S_{\circ})
 - (a) Micelle

(b) Multimolecular colloid

(c) Associated colloid

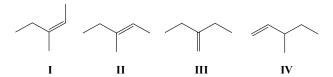
- (d) Macromolecular colloid
- Antacids are found in medicines that cure: **45.**
 - (a) Eye sight
- (b) Stomach ache (c) Pimples
- (d) Headache

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- 46. Which of the following compounds gives a primary alcohol upon reaction with phenylmagnesium bromide?
 - (a) 2-methyloxirane(b) Ethylene oxide
 - (c) Ethyl format

- (d) Carbon dioxide
- 47. What will be the product of the following reaction? $\frac{R_2NH}{R_2NH}$?
 - (a) 3-aminopyridine(b) 2-aminopyridine
 - (c) 3,5-diaminopyridine
- (d) 2,5-diaminopyridine
- 48. What will be the product of the following reaction? POCl₃
 4-pyridone
 - (a) 3-chloropyridine

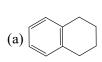
- (b) 2-dichloropyridine
- (c) 3,5–dichloropyridine
- (d) 2,5-dichloropyridine
- 49. Which of the following akenes yield(s) 3-bromo-3-methylpentane as the major product upon addition of HBr?

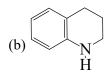


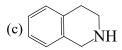
- (a) I and II only
- (b) III only
- (c) I, II and III only (d) All of them
- 50. What will be the atomic number of the element, if the last entering electron in an element has quantum number n = 3, l = 2, m = +2 and s = +1/2.
 - (a) 13
- (b) 21
- (c) 29
- (d) 39
- 51. What is the major organic product of the following reaction?

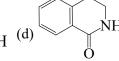
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- **52.** The least basic among the following is
 - $(a) Al(OH)_3$
- (b) La(OH)₃
- (c) Ce(OH)₃
- (d) Lu(OH)₃
- 53. Which one of the following compounds undergoes bromination of its aromatic ring (electrophilic aromatic substitution) at the fastest rate?

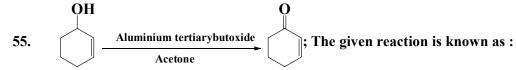








- 54. The pair of having similar geometry is:
 - (a) BF, and NH,
- (b) H₂O and C₂H₃ (c) CO₂ and SO₃
- (d) NH, and PH,



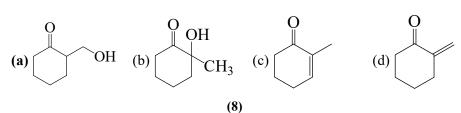
(a) Kolbe reaction

(b) Tischenko reaction

(c) MPV reaction

- (d) Oppeneur oxidation
- **56.** Which of the following statements is not true with respect to chemisorptions?
 - (a) Depends on nature of adsorbate and adsorbent
 - (b) Has a large heat of adsorption
 - (c) Forms a unimolecular layer
 - (d) Occurs at low temperature
- 57. What are the dispersed phase and dispersion medium in alcohol respectively?
 - (a) Alcohol, water (b) Solid, water
- (c) Water, alcohol (d) Solid, alcohol

PCC → (A) + HCHO $\xrightarrow{\text{H}_2\text{SO}_4}$ (B). Product B is : 58.



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- 59. Which of the following cannot be formed from Sandmeyer reaction on benzenediazonium chloride?

 - (a) Chlorobenzene (b) Bromobenzene (c) Iodobenzene
- (d) Benzonitrile
- 60. Which of the following can be produced by Gatterman reaction of diazonium salts?
 - (a) Bromobenzene (b) Fluorobenzene (c) Nitrobenzene

- (d) Cyanobenzene
- 61. The correct order of reactivity of p-halo nitrobenzene in the following reaction is

$$X = F, Cl, Br, 1$$

$$NO_2 \quad NaOMe \quad NO_2$$

$$MeO$$

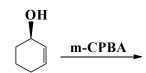
- (a) p-chloronitrobenzene > p-iodonitrobenzen > p-fluorontorbenzene > p-bromonitrobenzen
- (b) p-iodonitrobenzen > p-chloronitrobenzene > p-bromonitrobenzen > p-fluorontorbenzene
- (c) p-iodonitrobenzen > p-bromonitrobenzen > p-chloronitrobenzene > p-fluorontorbenzene
- (d) p-bromonitrobenzen > p-fluorontorbenzene > p-iodonitrobenzen > p-chloronitrobenzene
- **62.** Consider the following: According the Werner's theory
 - **A.** Ligands are connected to the metal ions by covalent bonds
 - **B.** Secondary valence is have directional properties
 - C. Secondary valences are non-ionisable of these statements choose the correct answer from the options given below:

Choose the correct option:

- (a) A, B and C only (b) B and C only
- (c) A and C only
- (d) A and B only
- **63.** According to MO theory, for the diatomic species, C,
 - (a) Bond order is zero and it is diamagnetic
 - (b) Bond order is two and it is paramagnetic
 - (c) Bond order is two and it is diamagnetic
 - (d) Bond order is zero and it is paramagnetic
- 64. The number of microstates present in ³F term is
 - (a) 3
- (b) 21
- (c) 9
- (d) 28
- How many M M bonds are present in [Cp Mo(CO)₃],? **65.**
 - (a) 1
- (b) 2
- (c) 0
- (d) 4

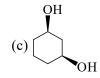
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66. When (R)-cyclohex-2-en-1-ol is treated with acid below gives









- (d) None of these
- 67. What will be the order of rotation barriers about C-N bond among these compounds?

(I)
$$CH_3$$
— CH_2 - NH_3 (II) C
 NH
 C
 NH
 H
 C
 NH_2

- (a) I > II > III
- (b) I < II < III
- (c) II > III > I
- (d) III > I > II
- 68. Which one of the following has the smallest heat of hydrogenation?
 - (a) 1-Butene
- (b) trans-2-Butene (c) cis-2-Butene
- (d) 1,3–Butadiene
- 69. The equivalent conductance of Ba²⁺ and Cl⁻ are respectively 127 and 76 ohm⁻¹ cm⁻¹ eq⁻ at infinite dilution. The equivalent conductance of BaCl, at infinite dilution will be:
 - (a) 101.5
- (b) 139.5
- (c) 203
- (d) 279

- 70. For spontaneous adsorption of gas:
 - (a) ΔS is positive so ΔH should be negative
 - (b) ΔS is negative so ΔH should be highly positive
 - (c) ΔS is negative so ΔH should be highly negative
 - (d) ΔS is positive so ΔH should also be highly positive
- 71. What is the catalyst used in manufacture of Ammonia by Haber's process?
 - (a) Fe
- (b) Molybdenum
- (c) Silica
- (d) Aluminium
- 72. The concentrations of a species A undergoing the reaction $A \rightarrow P$ is 1.0, 0.5, 0.33, 0.25 mol dm⁻³ at t = 0, 1, 2 and 3 seconds, respectively. The order of the reaction is
 - (a) 2
- (b) 1
- (c) 0
- (d) 3
- 73. A nucleotide is formed of which of the following units?
 - (a) Nitrogen base and phosphate
- (b) Nitrogen base, sugar and phosphate

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(c) Nitrogen base and sugar

(d) Sugar and phosphate

74. To which of the following does guanine form hydrogen bonds in DNA?

(a) Adenine

(b) Thymine

(c) Cytosine

(d) Guanine

75. Glycine and proline are the most abundant amino acids in which of the following structure?

(a) Hemoglobin (b) Myoglobin (c) Insulin

(d) Collagen

76. Which is not a type of catalyst?

(a) Positive catalyst

(b) Negative catalyst

(c) Auto catalyst

(d) Homogeneous catalysis

77. Mg_2C_3 reacts with water forming propyne, C_3^{4-} has:

(a) Two sigma and two pi bonds

(b) Three sigma and one pi bonds

(c) Two sigma and three pi bonds

(d) Two sigma and one pi bonds

78. In the formation of N_2 , from N_2 , the electron is removed from :

(a) σ orbital

(b) π orbital

(c) σ^* orbital

(d) π * orbital

79. XeF, is isostructural with

(a) ICl,

(b) SbCl,

(c) BaCl,

(d) TeF,

80. What is the reason for iodoform to be used as an antiseptic?

(a) Due to its unpleasant odour

(d) Due to its melting point

(c) Due to its solubility in alcohol

(d) Due to the liberation of free iodine

81. Which of the following complex has a highest oxidation state of metal?

(a) $(\eta^6 - C_6 H_6)_{2} Cr$ (b) $Mn(CO)_{5} Cl$

(c) $Na_2[Fe(CO)_4]$ (d) $K[Mn(CO)_5]$

82. Which among the following product is formed when ethyne undergoes hydrogenation?

(a) Formaldehyde (b) Formic acid

(c) Acetaldehyde

(d) Acetic acid

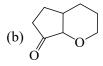
83. What is X in the following conversion?

$$\begin{array}{c}
OH \\
O \\
\end{array}$$

$$X$$

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84. Find the final product (C) in the following reaction is

$$OEt \xrightarrow{OH OH} (A) \xrightarrow{2 \text{ PhMgBr}} (B) \xrightarrow{H_3O^+} (C)$$

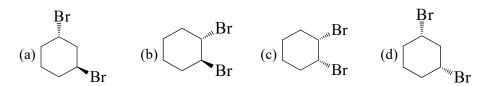
- 85. The number of substitution products formed when metabromo anisole is treated with KNH_2/NH_3 ?
 - (a) 1
- (b) 2
- (c) 3
- (d) 4
- 86. The equilibrium constant (K) of a redox reaction is related to the standard potential, E° , by the equation
 - (a) $ln(K) = -(nFE^{\circ}/RT)$
- (b) $ln(K) = (nFE^{\circ}/RT)$
- (c) $ln(K) = (RT/ nFE^{\circ})$
- (d) $ln(K) = -(RT/nFE^{\circ})$
- 87. At which of the following temperatures the reaction $Br_2(l) + Cl_2(g) \rightarrow 2BrCl(g)$ will be at equilibrium if $\Delta H = 30$ kJ mol⁻¹ and $\Delta S = 170$ J K⁻¹ mol⁻¹?
 - (a) 273 K
- (b) 285.7 K
- (c) 300 K
- (d) 450 K
- 88. Which of the following is an associated colloid?
 - (a) Starch
- (b) Soap
- (c) Proteins
- (d) Sol of gold
- 89. What will be the value of Kp if the Kc is 26 for a reaction, PCl₅

 → PCl₃ + Cl₂ at 25 degree Celsius:
 - (a) 0.46
- (b) 0.57
- (c) 0.61
- (d) 0.83

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- 90. When Q (concentration quotient) is smaller than equilibrium constant reaction will:
 - (a) Be in equilibrium

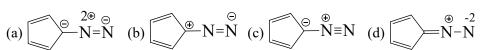
- (b) Proceed in forward direction
- (c) Proceed in backward direction
- (d) None of the above
- 91. Identify major product for the following reaction (i) HBr (ii) Br



- 92. Identify the correct relation between Δ_0 and Δ_t where Δ_0 denotes crystal field splitting in octahedral complexes and Δ_t denotes crystal field splitting in tetrahedral complexes
 - (a) $\Delta_{o} < \Delta_{t}$
- (b) $\Delta_0 > \Delta_1$
- (c) $\Delta_{\rm o} = \Delta_{\rm t}$
- (d) $\Delta_{o} \geq \Delta_{t}$
- 93. The complex [Co(H₂O)₆]³⁺ absorbs the wavelength of light corresponding to orange colour. Predict the colour of the coordination compound based on this information:
 - (a) Red
- (b) Yellow
- (c) Blue
- (d) Colourless
- 94. The absolute configuration at the two chiral cdntres of (–)-camphore is



- (a) 1R, 4R
- (b) 1R, 4S
- (c) 1S, 4R
- (d) 1S, 4S
- 95. The most stable resonating structure of the following compound is N = N



- 96. All the metals form oxides of the type MO except
 - (a) Copper
- (b) Barium
- (c) Silver
- (d) Lead

- 97. How many mmols of NaOH will be used in the titration with 33ml of 3 M HCl to form NaCl and water?
 - (a) 3 mmol
- (b) 10 mmol
- (c) 33 mmol
- (d) 100 mmol
- 98. In the given reaction the main product will be

- 99. Which one is the most stable carbonate?
 - (a) BaCO,
- (b) BeCO,
- (c) CaCO,
- (d) MgCO₃
- 100. Which of the ions have maximum hydration energy?
 - (a) Sr^{2+}
- (b) Ca²⁺
- (c) Mg^{2+}
- (d) Be^{2+}
- 101. Which of the following processes is used to do maximum work done on the ideal gas that is compressed to half of its initial volume?
 - (a) Adiabatic
- (b) Isobaric
- (c) Isothermal
- (d) Isochoric
- 102. K for a zero-order reaction is 2×10^{-2} mole/L/ sec. If the concentration of reactant after 25 sec is 0.5M, the initial concentration must have been:
 - (a) 0.5M
- (b) 0.75M
- (c) 1.0M
- (d) 1.25M

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103. CHO
$$\xrightarrow{\text{conc. KOH}} (A) \xrightarrow{H^{\oplus}} (B)$$
 cyclic product; Structure of (B) is:

(a)
$$COOH$$
 $COOK$ CH_2OH (c) $COOK$ CH_2OH

- 104. The type of viscosity which is related with the molecular weight of the polymer is:
 - (a) Coefficient of viscosity
- (b) Specific viscosity

(c) Reduced viscosity

(d) Intrinsic viscosity

	F	GT I GIC I KVS	I NVS I BPSC I	DSSSB ———
105.		ollowing complexes of (b) [Pt(bn) ₂] ²⁺		
106.	Which of the fo	ollowing is not a prot (b) PO(OH) ₃	tonic acid? (c) SO(OH) ₂	(d) SO ₂ (OH) ₂
107.	Increasing orders is:	er of stability among	the three main co	nformations of 2-fluoroethanol
	(a) Eclipse,gauc (c) Eclipse,anti,		(b) Gauche, eclip (d) Anti, gauche,	
108.	Permutit is: (a) A mixture of (c) Sodium alun	`aluminium oxide and	l silica	(b) Aluminosilicates (d) None
109.	Carboxy peption (a) Fe	lase is enzyme of wh (b) Mg	ich metal : (c) Cu	(d) Zn
110.	(a) Electron with(b) Electron rele	hdrawing groups easing groups n withdrawing and rel	C	her in the presence of :
111.	(a) A place when(b) A place when(c) An area belo	term "metastable" in re the composition of re the pressure remain w which a vapour—lie ere the temperature re	the system remains as constant with low quid mixture is obta	value
112.	eutectic temper (a) The system i (b) The system i (c) Components	rature?		stem that has reached the
113.	Which of the fo (a) It is a soapy (c) It is a dibasio	•	t sulphuric acid? (b) It is a monob (d) It is a tribasion	

(15)

114.	4. When an acid reacts with a metal, which one of the following gases is usually liberated?						
	(a) Ammonia gas	(b) Chlorine	(c) Oxygen	(d) Hydrogen gas			
115.	In the Lindeman mechanism of uninconcentration is:		olecuair reactions,	the observed order at low			
	(a) 0.5	(b) 1	(c) 1.5	(d) 2			
116.	According to Arrh (a) ln K decreases l (c) ln K increases l	linearly with 1/T	= rate constant and (b) ln K decreases (d) ln K increases l	-			
117.	Roots of Madar p	lant contains –					
	(a) Indigo	(b) Alizarin	(c) Saffron	(d) Crystal violet			
118.	The starting mate (a) Monochlorotrif (c) Vinyl chloride	rials of PTFE are: luoro ethylene	(b) Tetrafluoroethy(d) Styrene	lene			
119.	What is the major H ₂ SO ₄ at 413 K? (a) Ethene (c) Methoxyethane		(b) Methoxymethan (d) Ethoxyethane	drated with concentrated			
120.	substance disinteg	ife of a radioactive grates in 40 minutes (b) 15 min 20 sec	s?	of any given amount of the (d) 15 min 40 sec			
121.	$(_{5}I^{128}, t_{1/2} = 25 \text{ min})$) after 50 min. the i	remaining quantity	will be			
	(a) Half	(b) Third	(c) Quarter	(d) Nothing			
122.	Crystal field splitt	ting energy for octa	hedral $\left(\Delta_{_{f o}} ight)$ and te	trahedral $\left(\Delta_{_{\mathbf{t}}} \right)$ complexes is			
	(a) $\Delta_{\rm t} = \frac{4}{9} \Delta_{\rm o}$	(b) $\Delta_{\rm t} = \frac{1}{2} \Delta_{\rm o}$	(c) $\Delta_o = 2\Delta_t$	(d) $\Delta_{\rm o} = \frac{4}{9} \Delta_{\rm t}$			
123.	Which is the corre	ect order of ionic siz	zes? (At. no. : Ce =	58, Sn = 50,Yb = 70 and Lu			
	(a) Ce > Sn > Yb > (c) Sn > Ce > Yb >		(b) Sn > Yb > Ce > (d) Lu > Yb > Sn >				

= PGT | GIC | KVS | NVS | BPSC | DSSSB =

- 124. Which of the following shows disproportionation reactions?
 - (a) $[Au(H_2O)_6]^{2+}$
- (b) $[Ni(H_2O)_6]^{2+}$
- (c) $[Cu(H_2O)_6]^{2+}$
- (d) $[Au(CN)_4]^{3-}$
- 125. Incorrect statement regarding pairing energy is/are
 - (a) Pairing energy decrease for heavier congeners
 - (b) Covalency in metal ligand bond decrease with pairing energy
 - (c) pi acid ligand can decrease the pairing energy
 - (d) Pairing energy decrease with increase in positive oxidation state of metal

	ANSWER KEY												
1	(b)	11 (b)	21 (a)	31 (a)	41 (d)	51 (a)	61 (c)	71 (a)	81 (c)	91 (a)	101(a)	111(d)	121(c)
2	(b)	12 (a)	22 (b)	32 (b)	42 (a)	52 (a)	62 (b)	72 (a)	82 (c)	92 (b)	102(c)	112(a)	122(a)
3	(b)	13 (d)	23 (c)	33 (c)	43 (c)	53 (b)	63 (c)	73 (b)	83 (c)	93 (c)	103(a)	113(c)	123(a)
4	(a)	14 (b)	24 (c)	34 (a)	44 (b)	54 (d)	64 (b)	74 (c)	84 (b)	94 (c)	104(d)	114(d)	124(a)
5	(d)	15 (a)	25 (a)	35 (a)	45 (b)	55 (d)	65 (a)	75 (d)	85 (c)	95 (c)	105(d)	115(d)	125(b)
6	(c)	16 (b)	26 (a)	36 (b)	46 (b)	56 (d)	66 (b)	76 (d)	86 (b)	96 (c)	106(a)	116(a)	
7	(b)	17 (a)	27 (d)	37 (c)	47 (b)	57 (d)	67 (b)	77 (a)	87 (b)	97 (d)	107(c)	117(b)	
8	(d)	18 (b)	28 (d)	38 (d)	48 (b)	58 (a)	68 (d)	78 (a)	88 (b)	98 (a)	108(b)	118(a)	
9	(d)	19 (c)	29 (d)	39 (c)	49 (d)	59 (c)	69 (d)	79 (a)	89 (c)	99 (a)	109(d)	119(d)	
10	(d)	20 (d)	30 (d)	40 (b)	50 (b)	60 (a)	70 (c)	80 (d)	90 (b)	100(d)	110(a)	120(a)	

Solution

- (b) In isothermal process →
 Entropy increase enthalpy remains constant
- 2. **(b)** $w = -P_{ext.} \Delta V$ = -1 × (10 - 1) W = -9 L-bar $\therefore 1L - bar = 100 J$

$$\omega = -9 \times 100 \text{ J}$$

work done $\omega = -0.9 \text{ kJ}$

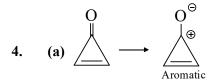
3. **(b)** r = 0.14 nmfor FCC $4r = a\sqrt{2}$ $r \rightarrow \text{ atomic radius}$ $a \rightarrow \text{ length of a side of cell}$

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$$r=\frac{a}{2\sqrt{2}}$$

$$0.14 \text{ nm} = \frac{a}{2 \times 1.414}$$

a = 0.4 nm



Due to aromatic nature above compound shows maximum dipole moment.

5. (d)
$$\xrightarrow{\text{COO}^{\Theta} \text{Ag}^{\oplus}} \xrightarrow{\text{COO}^{\Theta} \text{Ag}^{\oplus$$

$$\begin{array}{c} \triangle \\ \hline \text{decarboxylatia} \\ \\ \downarrow \\ \text{COOH} \\ + \text{CO}_2 \\ \end{array}$$
Nicotinic acid

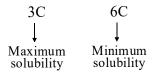
7. (b) Exter range of IR → 1750 – 1735 cm⁻¹
 In cyclic esters C = O frequency increases with decreasing ring size.

3. (d)
$$\longrightarrow$$
 \longrightarrow $+$ Br $\xrightarrow{\text{AgNO}_3}$ AgBr

As the length of hydrophobic part increases. Solubility in water decreases.

(18)

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10. (d) Witting reaction

$$O \xrightarrow{Ph_3P = CH_2} CH_2$$

Mechanism

$$\begin{array}{c} & \overset{\text{\tiny \textcircled{e}PPh_3}}{\bigcap} \\ & \overset{\text{\tiny \textcircled{e}CH_2}}{\bigcap} \\ & & \overset{\text{\tiny \textcircled{e}PPh_3}}{\bigcap} \\ & & & \overset{\text{\tiny \textcircled{e}CH_2}}{\bigcap} \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & \\ & & \\ & & & \\ & & \\ & & \\ & & & \\ & & \\ & & & \\ & & \\ & & \\ & & \\ & & & \\ & & \\ &$$

- 11. **(b)**
- **12. (a)** A necessary condition for osmosis is semi-permeable membrane.
- 13. (d) Most all the methods given in the option are used to determine the molar mass of bimolecular but osmosis method is most appropriate and most widely used.

14. **(b)**
$$t_{1/2} = \frac{2^{n-1}}{n-1} \times \frac{1}{K.A_0^{n-1}}$$

$$t_{1/2} \alpha \frac{1}{A_0^{A-1}}$$

Hence ty₂ for nth hat order is inversity proportional to (n-1)th power of initial concentration.

15. (a)
$$K = \frac{2.303}{t} \log \frac{A_o}{A}$$

$$K = \frac{2.303}{20} \log \frac{100}{90}$$

$$K = \frac{2.303}{20} (\log 10 - \log 9)$$

$$K = \frac{2.303}{20} \times 0.0458$$

$$t = \frac{2.303}{20} \log 10 \frac{A_o}{A}$$

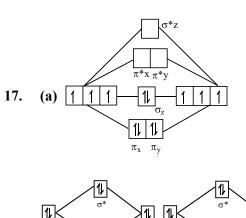
$$= \frac{2.303 \times 20}{2.303 \times 0.0458} \log \frac{100}{81} \times 100$$

$$= \frac{20}{0.0458} (\log 100 - \log 81)$$

$$= \frac{20}{0.0458} \times 0.915 = 39.96$$

$$t = 40 \min$$

16. (b)



Molecular orbital diagram of N₂

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Bond order =
$$\frac{\text{Bonding e}^- - \text{antibonding e}^-}{2}$$

Bond order = of
$$N_2 = \frac{6-0}{2} = 3$$

of
$$N_2^+ = \frac{5-0}{2} = 2.5$$

of
$$N_2^+ = \frac{6-1}{2} = 2.5$$

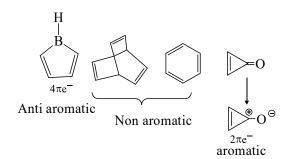
- **18. (b)** Electronic transitions are studied by UV spectrometer
- 19. (c)
- **20. (d)** Since Br is a very bulky group therefore stable is anti conformation.

$$H \xrightarrow{Br} H$$

Note

$$\begin{array}{c} Br \\ H \\ H \\ H \\ Br \\ Br \end{array}$$

21. (a) $4n \pi e^- \rightarrow Antiaromatic$ $(4n + 2) \pi e^- \rightarrow Aromatic$ Antiaromatic and aromatic compounds are planar, cyclic and have complete conjugation



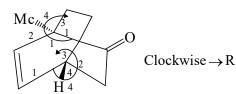
- 22. (c) $B \stackrel{\text{(..)}}{|} B$ lone pair = 1 B Bond pair = 3
- (c) complexes in which a metal is bound to only one kind of donar groups, eg → [Fe(CN)₆]^{3- Homoleptic complex}
 When a metal is bound more than one kind of donar groups, is called hetero lept complexes.

Eg:-
$$\left[\operatorname{Co}\left(\operatorname{NH}_{3}\right)_{4}\operatorname{Cl}_{2}\right]^{\oplus}$$

- 24. (c)
- 25. (a)
- **26.** (a) When lone pair of e^- of CO donate to vacant orbital of metal, σ bond forms.

When metal donate it's electrons to CO then π bond forms. Which is called back bonding.

- 27. (d) Octahedral geometry can have two type of hybridization.i.e. sp³d², d²sp³
- **28. (d)** To find out the configuration following steps should be followed
 - (i) Give priority to all 4 group (Higher mass have higher priority)
 - (ii) Rotate priority 1 to 4
 If clockwise rotation → R
 configuration
 If anticlockwise rotation → S
 configuration
 - (iii) If uth group is an wedge (-) Configuration will be change.



Anti clockwise \rightarrow S But 4th on wedge S \rightarrow R 1R, 4R

- 29. (d)
- 30. (d) Up to down → Atomic radius increase

 Left to right → Atomic radius decrease

 In isoelectronic species, higher the negative charge, higher the ionic radius and higher the positive charge, lower the ionic radius.

$$N^{3-} > O^{2-} > F^- > Ne > Na^+ > Mg^{++} > Al^{+++}$$

Hence option (d) is incorrect.

- 31. (a) On dilution \rightarrow
 - Molar conductance increase
 - Equivalent conductance increase
 - Specific conductance decrease
- 32. (b) Spectrochemical series \rightarrow

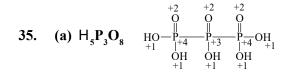
$$\begin{split} &CO>CN^{-}>en>NH_{_{3}}>EDTA^{^{4-}}>\\ &NCs^{-}>H_{_{2}}O>O^{^{-2}}>C_{_{2}}O_{_{4}}^{^{2-}}>OH^{^{0}}\\ &>F^{^{\oplus}}>Cl^{-}>SCN^{^{0}}>S^{-}>Br^{^{0}}>l^{^{0}} \end{split}$$

All the compounds given is options are in d^6 state. Hence Δ_o should be equal to all but strong field ligands (CO, CN $^-$, en etc.) have high Δ_o then low field ligands (I^0 , Br^0 , S^- etc.)

 $\Delta_{o} \text{ order} \rightarrow [\text{Co(CN)}_{6}]^{3-} > \\
[\text{Co(NH}_{3})_{6}]^{3+} > [\text{Co(C}_{2}\text{O}_{4})_{3}]^{3-} > \\
[\text{Co(H}_{2}\text{O})_{6}]^{3+}$

- 33. (c) On increasing the concentration of reactant in a reversible reaction, equilibrium concentrations will unchange.
- 34. (a) Super closo $[B_{n}H_{n}]^{2+}$ Hyper closo $[B_nH_n]^{\circ}$ Close $[B_{n}H_{n}]^{2}$ Nido $[B_{n}H_{n}]^{4-}$ Arachno $[B_n H_n]^{6-}$ [B_H_]8-Hypo $[B_{n}H_{n}]^{10}$ " Clado $[B_{12}H_{12}]^{2-} \rightarrow [B_nH_n]^{2-}$ closo structure

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- 36. (b) for combustion reaction $\Delta H = \sum_{\text{Reactant}} \sum_{\text{Product}} H_{\text{Product}}$
- 37. (c) → optically inactive

 2-Bromo propane

 → optically inactive

 1-Bromo 3 methyl Butane

 H

 * optically active → Exists as a pair of enantiomers Cyclo hex-2-

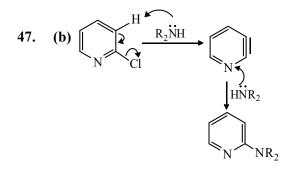
ene-1-ol

- Note→ Compound having plane of symmetry (POS) are optically inactive and achiral.
- **39. (c)** <u>Chiral carbon</u> → An carbon atom that is bonded to four different types of atoms or groups or groups of atoms.

Hence 3 chiral atoms are present.

- 40. (b) Both compounds are identical
- 41. (d) At NTP At STP
 Volume = 22.4 L
 Temp = 298 K
 Pressure = 1 atm 1 atm
- 42. (a) Average speed $V_{avg} = \sqrt{\frac{8RT}{\pi M}}$ $V_{ang} \alpha \sqrt{\frac{T}{M}} = (V_{avg})O_2 = (V_{av})CH_4$ $\sqrt{\frac{T_{O_2}}{M_{O_2}}} = \sqrt{\frac{T_{CH_4}}{M_{CH_4}}}$ $\sqrt{\frac{300}{32}} = \sqrt{\frac{T_{CH_4}}{16}}$ $T_{CH_4} = 300K$
- 43. (c) H_3^{\oplus} $H_3^{-60^{\circ}}$
- 44. (b) Associated colloid → Micelles, soap
 Macro molecular colloid → Starch,
 Cellulose, Proten and enzymes.
 Multimolecular colloid → Gold, sol, S₈
- **45. (b) Antacides cures** → Stomach ache

PGT | GIC | KVS | NVS | BPSC | DSSSB

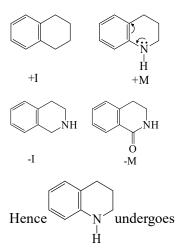


- 48. (b)
- 49. (d)
- 50. **(b)** n = 3 1 = 2 (d subshell) m = +2 s = +1/2 $3d^{1}$ \downarrow n = 3, 1 = 2, m = +2, s = +1/2

Atomic number (z) = 21 Electronic configuration \rightarrow [Ar]4s²3d¹

51. (a)
$$\underbrace{ (1) \, k M_n O_4, OH}^{\Theta}, Cold \longrightarrow OH$$

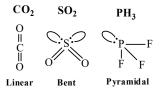
- 52. (a) Al(OH), are lewis acid, rest are base.
- **53. (b)** Electron donating group increase the rate of the electrophilic aromatic substitution and vice versa.



bromination with fastest rate.

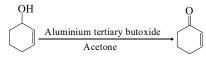
54. (d) BF₃ NH₃ H₂O

Trigonal Pyramidal Bent



Hence NH₃ and PH₃ have similar geometry.

55. (d) Oppeneur oxidation



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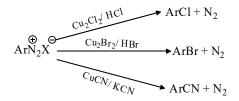
- **56. (d)** Characteristics of chemisorptions.
 - i. Chemical bond formation
 - ii. Highly specific in nature
 - iii. Irreversible
 - iv. Enthalpy of adsorption is high (80 240 kJ mol)
 - v. High temperature is favourbles
 - **vi.** High activation energy is required.
 - vii. Unimolecular layer forms
 - viii.Depends on nature of adsorbate and adsorbent.

Hence option (d) is correct.

57. (d) Colloidal Dispersion Dispersd

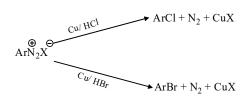
<u>solution</u>	<u>medium</u>	<u>phase</u>
Aerosol	Gas	Solid/liq.
Hydrosol	water	Solid
Alcohol	Alcohol	Solid

- 58. (a)
- 59. (c) Sandmayer reaction \rightarrow



<u>Note</u> – Iodobenzene can't be formed by sandmeyer reaction.

60. (a) <u>Gatterman reaction</u> \rightarrow



61. (c) Rate I > Br > Cl > F

- **62. (b)** Werner's theory of co-ordination 1892 Compounds → main postulates are
 - Two types of valenes → primary & secondary
 - **2)** Primary valences are ionisable and non-directional, satisfied by negative ions.
 - 3) Secondary valence are non-ionisable and directional, satisfied by neutral molecules or negative ions.
 - 4) Primary valences showed by dotted line (.....) while secondary valences showed by thick line (——)

63.	(c) C ₂ =	12e	Bond order = $\frac{4-0}{2}$ = 2
		1, 1, 8e	zero unpaired electron lence diamagnetic

64. (b) Micro states = (2S + 1)(2L + 1)³F L = 3

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$$2S + 1 = 3$$

Micro states = $3 \times (2 \times 3 + 1)$
= $3 \times 7 = 21$

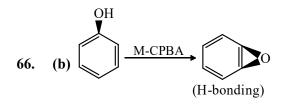
65. (a)
$$[C_p Mo(CO)_3]_2$$

 $VE = 2[5 + 6 + 2 \times 3] = 34$
 $18n - VE$

M - M bond = $\frac{18n - VE}{2}$ n = number of metals

$$=\frac{18\times2-34}{2}$$

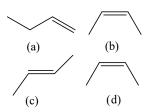
M-M bond = 1



67. (b) Rotation barrier α Bond strength(I) CH₂-CH₂-NH₂

C-N bond strength III > II > I Rotation Barrier III > II > I

68. (d) Heat of hydrogenation $\alpha \frac{1}{\text{stability}}$



Stability order d > c > b > aHeat of hydrogenation a > b > c > d(smallest HOH)

69. (d)
$$(\lambda_{eq}) Ba^{2+} = 127 \Omega^{-1} cm^{-1} eq^{-1}$$

 $(\lambda_{eq}) Cl^{-} = 76 \Omega^{-1} cm^{-1} eq^{-1}$
 $\lambda_{BaCl_{2}} = \lambda_{Ba^{2+}} + 2\lambda_{Cl^{-}}$
 $= 127 + 2 \times 76$
 $= 279 \Omega^{-1} cm^{-1} eq^{-1}$

- 70. (c) For spontaneous adsorption of gas
 - (1) Entropy change is positive
 - (2) ΔH should be highly negative
- 71. (a) Haber's process \rightarrow $N_2 + 3H_2 \xrightarrow{\text{Fe, Mo}} 2NH_3$ Fe \rightarrow Act as catalyst
 Mo \rightarrow Act as promoter
- 72. (a) For 2nd order reaction $\frac{1}{A_2} \frac{1}{A_1} = K(t_2 t_1)$ $A_2 = 0.5, A_1 = 1$ $t_2 = 1, t_1 = 0$ $\frac{1}{0.5} \frac{1}{1} = K(1 0)$

2 - 1 = K

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$$K = 1$$

$$A_{2} = 0.33, A_{1} = 0.5$$

$$t_{2} = 2, t_{1} = 1$$

$$\frac{1}{0.33} - \frac{1}{0.5} = K(2-1)$$

$$K = 1$$

$$A_{2} = 0.25, A_{1} = 0.33$$

$$t_{2} = 3, t_{2} = 2$$

$$\frac{1}{0.25} - \frac{1}{0.33} = K(3-2)$$

$$K = 1$$

Hence reaction is 2nd order reaction

Note → Fro 0th order reaction

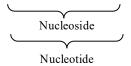
$$\frac{A_2 - A_1}{t_1 - t_2} = \frac{A_3 - A_2}{t_2 - t_3} = \frac{A_4 - A_3}{t_3 - t_4} = K$$

For 1st order reaction

$$\frac{\log_{e} A_{2} - \log_{e} A_{1}}{t_{2} - t_{1}}$$

$$= \frac{\log_{e} A_{3} - \log_{e} A_{2}}{t_{3} - t_{1}} = -K$$

73. **(b)** Nitrogen base + sugar + Phosphate



- 74. (c) Hydrogen bonding in DNA
 C ≡ G (Cytosine and guanine)
 A=T (Adenine and thyning)
- 75. (d) Glycene, proline and hydroxyl proline contribute to 57% of total amino acids in collagen, which

accounts for one-third of proteins in animals.

76. (d)

77. (a)
$$C_3^{4}$$
 $^{-2}C\frac{\pi}{\sigma}C\frac{\pi}{\sigma}C^{-2}$

$$\sigma \text{ bond} = 2$$

$$\pi \text{ bond} = 2$$

For the formation of N_2^{\oplus} , electron will be removed from σ orbital.

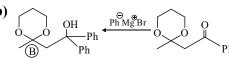
79. (a)
$$ICl_2^{\Theta}$$
 $Cl \xrightarrow{I}_{\Theta}$ Cl

$$XeF_2 \quad F \xrightarrow{Xe}_{G}$$

XeF_2	ICl_2^{Θ}	
Hybridizatio	on sp³d	sp^3d
Geometry	TBP	TBP
Shape	lnear	linear

80. (d) Iodoform is used as an antiseptic due to liberation of free iodine.

PGT | GIC | KVS | NVS | BPSC | DSSSB



85. (c)
$$\Delta G = \Delta G^{\circ} + RT \ln k$$

At equilibrium "G = 0

$$0 = \Delta G^{\circ} + RT \ln k$$

$\Delta G^{\circ} = -RT \ln k$

$$\ln k = \frac{\Delta G}{-RT} :: \Delta G^{\circ} = -nFE^{\circ} \text{ cell}$$

$$\ln k = \frac{-nFE_{cell}^{\circ}}{-RT}$$

$$\ln k = \frac{nF E_{cell}^{o}}{RT}$$

90. **(b)**
$$Q_c = K_c$$
 equilibrium $Q_c < K_c$ forward direction $Q_c > K_c$ Backward direction $Q_c \rightarrow$ concentration quotient, $K_c \rightarrow$ equilibrium constant

92. **(b)**
$$\Delta t = \frac{4}{3} \Delta_o$$

 $\Delta_{sp} = 1.3 \Delta_o$

$$\Delta_{\rm sp} = 1.3\Delta_{\rm o}$$

$$\Delta_{\rm sp} > \Delta_{\rm o} > \Delta_{\rm t}$$

tetrahedral planar

93. (c) When a compound absorbed any particular wavelength of a particular Then it shows, colour. complementary colour.

Colour cycle of complementary <u>colour</u>



If compound absorbed

orange coloure, the colours of compound will be Blue.

94. (d) Anticlock wise \rightarrow S



wedge $R \rightarrow S$ 1S, 4S

THE RASAYANAM

95. (c)
$$N = N \oplus N \oplus N = N$$

- 96. (c) MO type \rightarrow CuO, BaO, PbO
- (d) 33 ml of 3 M HCl 97.

$$M = \frac{n}{V} \times 1000$$

$$3 = \frac{n_{HCl}}{33} \times 1000$$

$$\begin{aligned} n_{HCl}^{} &= 100 \times 10^{\text{-3}} \text{ mol} \\ n_{HCl}^{} &= 100 \text{ m mol} \end{aligned}$$

Hence 100 m mol NaOH required

98. (a)

99. (a) Thermal stability of carbonates of alkaline earth metal increases down the group.

$$BeCO_3 < MgCO_3 < CaCO_{3-} < SrCO_3 < BaCO_3$$

100. (d) Hydration energy $\propto \frac{\text{Charge}}{\text{size}}$

Size
$$Be^{2+} < Mg^{+2} < Ca^{+2} < Sr^{+2} < Ba^{++}$$

 $\label{eq:hydration} \textbf{Hydration} \ \ Be^{\text{+2}} \! > \! Mg^{\text{+2}} \! < \! Ca^{\text{+2}} \! > \! Sr^{\text{+2}} \! > \! Ba^{\text{++}}$ energy

101. (b) Work done

Isobaric > Isothemal > Adiabatic > Isochoric

102. (c) $K = 2 \times 10^{-2} \text{ mol L}^{-1} \text{ sec}^{-1}$ t = 25 sec

$$t = 25 \text{ sec}$$

 $A = 0.5 \text{ M}$

$$A = A - Kt$$

$$A = A_0 - Kt$$

$$A = A_o - Kt$$

 $0.5 = A_o - 2 \times 10^{-2} \times 25$

$$A_0 = 0.5^{\circ} +$$

$$A_0 = 0.5 + 0.5$$

$$A_0 = 1M$$

103. (a)

104. (d) Mark – Kuhn – Houw ink equation \rightarrow

$$[n] = K \left(\overline{M}_{v}\right) \alpha$$

 $[n] \rightarrow Intrinsic viscosity$

K, $\alpha \rightarrow constant$

 $M_{y} \rightarrow Molecular mass$

105. (d)
$$e^{n}$$
 Pros \rightarrow

Optically inactive

Protic acid

Protic acid

= PGT | GIC | KVS | NVS | BPSC | DSSSB :

Lewis acid

Protic acid

Stability Gauche > Anti > Eclipse

- **108. (b)** Per mutit is → Aluminosilicates
- 109. (d) Carboxy peptidase

110. (a) In the presence of electron electron withdrawing group rate of nucleophilic substitution reactions increases

$$\begin{array}{c|c}
Cl & Cl \\
\hline
NaOH \\
O_2N & NO_2
\end{array}$$

$$\begin{array}{c|c}
Cl & NO_2 & Cl \\
NO_2 & NO_2
\end{array}$$

$$\begin{array}{c|c}
NO_2 & NO_2
\end{array}$$

$$\begin{array}{c|c}
NO_2 & NO_2
\end{array}$$

111. (d)

- 112. (a)
- 113. (c) Sulphuric acid is a dibasic acid

- 114. (d) $M + HCl \rightarrow MCl + H_2 \uparrow$
- 115. (d) Lindemann equation

Rate =
$$\frac{dP}{dt} = \frac{K_1 K_2 [A^2]}{K_{-1}[A] + K_2}$$

At law concentration $K_1[A] \le K_2$

$$\frac{dP}{dt} = \frac{K_1 K_2 \left[A^2\right]}{K_2 + K_2}$$

$$\frac{dP}{dt} = \frac{K_1}{2} \left[A^2 \right]$$

2nd order kinetic

116. (a)
$$\left[K = Ae^{-Ea/RT}\right]$$

$$lnk = ln A - \frac{Ea}{RT}$$

$$y = c + m x$$



Hence ln K decreases lineary with time

117. (b) Roots of modar plant contains Alizarin.

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$$F_2C=CF_2$$

118. (b) Tetrafluoro ethylene

Polymerisation
$$\frac{\left(F_2C - CF_2\right)_n}{\left(PTFE\right)}$$
Polytetra fluoro ethylene (PTFE)

119. (d) Mechanism

(I)
$$CH_3-CH_2-\ddot{\square}-H+H \xrightarrow{\oplus} CH_3-CH_2-\overset{H}{\overset{}{\circ}}-H$$

120. (a)
$$3 \times t_{1/2} = t_{87.5\%}$$

 $3 \times t_{1/2} = 40 \text{ min}$
 $t_{1/2} = 13 \frac{1}{3} \text{min}$
 $t_{1/2} = 13 \text{ min } 20 \text{ sec}$

121. (c)
$$t_{1/2} = 25 \text{ min}$$

$$K = \frac{0.693}{t_{1/2}}$$

$$K = \frac{0.693}{25} = \frac{2.303 \times \log 2}{25}$$

$$t = \frac{2.303}{K} \log \frac{A_o}{A}$$

$$56 = \frac{2.303 \times 25}{2.303 \times \log 2} \times \log \frac{A_o}{A}$$
$$2 \times \log 2 = \log \frac{A_o}{A}$$
$$\log 2^2 = \log \frac{A_o}{A}$$
$$4 = \frac{A_o}{A}$$
$$A = \frac{A_o}{A}$$

- 122. (a)
- 123. (a) Ce, Xb & Lu are uf series elements and in uf series on going left to right ionic radius decreases.

- 124. (a)
- 125. (b)